Pharmacology A Patient Centered Nursing Process Approach 8th Edition Kee Test Bank

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Chapter 02: The Drug Approval Process Test Bank

MULTIPLE CHOICE

- The nurse is preparing to administer a schedule II injectable drug and is drawing up half of the contents of a single-use vial. Which nursing action is correct?
 - Ask another nurse to observe and cosign wasting the remaining drug from the vial.
 - b. Keep the remaining amount in the patient's drawer to give at the next dose.
 - Record the amount unused in the patient's medication record.
 - Dispose of the vial with the remaining drug into a locked collection box.

ACMEST AND

Schedule II drugs are controlled substances, and all must be accounted for. When wasting a portion of a drug, another nurse should observe and cosign that a drug was wasted.

DRF: COGNITIVE LEVEL: Applying (Application) REF: Page 16

TOP: NURSING PROCESS: Nursing Intervention

MSC: NCLEX: Physiological Integrity: Pharmacological and Parenteral Therapies

- A patient is prescribed a medication and asks the nurse if the drug is available in a generic form. The nurse understands that a generic medication will have a name that
 - a. is a registered trademark.
 - b. is always capitalized.
 - describes the drug's chemical structure.
 - is non-proprietary.

ANS: ID

The generic name is the official, non-proprietary name for a drug. The brand name is the trademark name and is always capitalized. The chemical name describes the chemical structure of the drug.

DIF: COGNITIVE LEVEL: Understanding (Comprehension) REF: Page 18

TOP: NURSING PROCESS: N/A

MSC: NCLEX: Physiological Integrity: Pharmacological and Parenteral Therapies

- 3. A patient receives a prescription on which the provider has noted that a generic medication may be given. The patient asks the nurse what this means. What will the nurse tell the patient about generic drugs?
 - They contain the same inert ingredients as brand-name drugs.
 - They have chemical structures that are identical to proprietary drugs.
 - c. They tend to be less expensive than brand-name drugs.
 - d. They undergo extensive testing before they are marketed.

ANS: C

Generic drugs are approved by the FDA if they are proved to be bioequivalent to the brand-name drug. They tend to be less expensive because manufacturers of these drugs do not have to do the extensive testing required of brand-name drugs before marketing. They are not identical to brand-name drugs and often have different mert ingredients.

DIF: COGNITIVE LEVEL: Applying (Application) REF: Page 18

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Table of Contents Kee Pharmacology Chapter 1test Bank

- 1. Understanding the eBook Kee Pharmacology Chapter 1test Bank
 - The Rise of Digital Reading Kee Pharmacology Chapter 1test Bank
 - Advantages of eBooks Over Traditional Books
- 2. Identifying Kee Pharmacology Chapter 1test Bank
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
- 3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Kee Pharmacology Chapter 1test Bank
 - User-Friendly Interface
- 4. Exploring eBook Recommendations from Kee Pharmacology Chapter 1test Bank
 - Personalized Recommendations
 - Kee Pharmacology Chapter 1test Bank User Reviews and Ratings
 - Kee Pharmacology Chapter 1test Bank and Bestseller Lists
- 5. Accessing Kee Pharmacology Chapter 1test Bank Free and Paid eBooks
 - Kee Pharmacology Chapter 1test Bank Public Domain eBooks
 - Kee Pharmacology Chapter 1test Bank eBook Subscription Services
 - Kee Pharmacology Chapter 1test Bank Budget-Friendly Options

- 6. Navigating Kee Pharmacology Chapter 1test Bank eBook Formats
 - o ePub, PDF, MOBI, and More
 - Kee Pharmacology Chapter 1test Bank Compatibility with Devices
 - Kee Pharmacology Chapter 1test Bank Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Kee Pharmacology Chapter 1test Bank
 - Highlighting and Note-Taking Kee Pharmacology Chapter 1test Bank
 - Interactive Elements Kee Pharmacology Chapter 1test Bank
- 8. Staying Engaged with Kee Pharmacology Chapter 1test Bank
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Kee Pharmacology Chapter 1test Bank
- 9. Balancing eBooks and Physical Books Kee Pharmacology Chapter 1test Bank
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Kee Pharmacology Chapter 1test Bank
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Kee Pharmacology Chapter 1test Bank
 - Setting Reading Goals Kee Pharmacology Chapter 1test Bank
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Kee Pharmacology Chapter 1test Bank
 - Fact-Checking eBook Content of Kee Pharmacology Chapter 1test Bank
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
- 14. Embracing eBook Trends
 - Integration of Multimedia Elements

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